

List of Clinical Tests by Disease

September 1, 2010

This list is for information only. It does not have to be returned with the clinical specimen.

Test No.	Test Name	Price	Test No.	Test Name	Price
100	Single exon sequencing in any gene that we test	\$ 190		Brachyactyly	
200	Double exon sequencing in any gene that we test	\$ 340	136	<i>ROR2</i> Sequencing – Tier 1 only	\$ 740
300	Triple exon sequencing in any gene that we test	\$ 390			
900	Non-Standard Test			Cardiac Channelopathy	
			492	<i>SCN5A</i> Sequencing	\$1,460
	Alagille Syndrome				
427	<i>JAG1</i> Sequencing	\$1,090		Cardio-Facio-Cutaneous (CFC) Syndrome	
			110	<i>BRAF, MAP2K1, MAP2K2, KRAS</i> Sequencing	\$490-990
	Amyotrophic Lateral Sclerosis		111	<i>BRAF</i> Sequencing – Tier 1 only	\$ 490
155	<i>ALS</i> Gene Sequencing Panel	\$1,440	112	<i>MAP2K1, MAP2K2, KRAS</i> Sequencing–Tier 2 only	\$ 590
108	<i>ALS2</i> Sequencing	\$1,790	113	<i>BRAF</i> Sequencing (remaining exons) – Test 2	\$ 790
154	<i>ANG</i> Sequencing	\$ 340	378	<i>KRAS</i> Sequencing	\$ 440
152	<i>FUS</i> Sequencing	\$ 890			
109	<i>SETX</i> Sequencing	\$1,690			
106	<i>SOD1</i> Sequencing	\$ 490		Cerebral Cavernous Malformations	
153	<i>TARDPB</i> Sequencing	\$ 470	120	Sequential <i>CCM</i> Gene Testing	\$190-1690
107	<i>VAPB</i> Sequencing	\$ 490	121	<i>KRIT1/CCM1</i> Sequencing	\$ 840
			122	<i>CCM2</i> Sequencing	\$ 740
600	Array CGH	See description on website	123	<i>PDCD10/CCM3</i> Sequencing	\$ 540
			124	<i>CCM2</i> Deletion Testing	\$ 190
			125	<i>KRIT1/CCM1</i> Common Hispanic Mutation	\$ 190
	Arrhythmogenic Right Ventricular Cardiomyopathy				
207	Sequential Testing of Four Desmosomal Genes	\$820-3810		CHARGE Syndrome	
206	<i>DSC2</i> Sequencing	\$ 900	130	<i>CHD7</i> Sequencing	\$1,890
205	<i>DSG2</i> Sequencing	\$ 940			
203	<i>DSP</i> Sequencing	\$1,820			
204	<i>PKP2</i> Sequencing	\$ 820		Classic Lissencephaly	
			503	<i>DCX</i> Sequencing	\$ 590
	Arthrogyrosis-Renal Dysfunction-Cholestasis (ARC) Syndrome		507	<i>PAFAH1B1/LIS1</i> Sequencing	\$ 670
591	<i>VIPAR</i> Sequencing	\$1,020		Congenital Contractural Arachnodactyly	
248	<i>VPS33B</i> Sequencing	\$1,090	395	<i>FBN2</i> Sequencing	\$2,190
	Autosomal Recessive Polycystic Kidney Disease				
101	<i>PKHD1</i> Sequencing	\$1,190-2990		Congenital Disorders of Glycosylation	
102	<i>PKHD1</i> Sequencing – Tier 1 only	\$1,190	540	Congenital Disorders of Glycosylation – Panel 1	\$490-1,470
103	<i>PKHD1</i> Sequencing – Tier 2 only	\$2,300	542	Congenital Disorders of Glycosylation – Panel 2	\$470-3,020
	Bardet-Biedl Syndrome		536	<i>ALG2</i> Gene Sequencing	\$ 470
269	Bardet-Biedl Syndrome Sequencing Panel	\$490-6,995	539	<i>ALG3</i> Gene Sequencing	\$ 590
254	<i>ARL6/BBS3</i> Sequencing	\$ 540	533	<i>ALG6</i> Gene Sequencing	\$ 740
252	<i>BBS1</i> Sequencing	\$ 890	537	<i>ALG8</i> Gene Sequencing	\$ 790
253	<i>BBS2</i> Sequencing	\$ 890	534	<i>ALG12</i> Gene Sequencing	\$ 590
255	<i>BBS4</i> Sequencing	\$ 890	535	<i>DPM1</i> Gene Sequencing	\$ 570
256	<i>BBS5</i> Sequencing	\$ 740	543	<i>DPM3</i> Gene Sequencing	\$ 340
258	<i>BBS7</i> Sequencing	\$1,020	531	<i>PMM2</i> Gene Sequencing	\$ 490
261	<i>BBS9</i> Sequencing	\$1,130	538	<i>MPDU1</i> Gene Sequencing	\$ 540
262	<i>BBS10</i> Sequencing	\$ 490	532	<i>MPI</i> Gene Sequencing	\$ 490
264	<i>BBS12</i> Sequencing	\$ 490			
257	<i>MKKS/BBS6</i> Sequencing	\$ 580		Congenital Generalized Lipodystrophy	
263	<i>TRIM32/BBS11</i> Sequencing	\$ 490	462	<i>AGPAT2</i> Sequencing	\$ 640
259	<i>TTC8/BBS8</i> Sequencing	\$ 850			

Test No.	Test Name	Price	Test No.	Test Name	Price
	Congenital Myasthenic Syndrome		664	<i>IMPDH1</i> Sequencing	\$ 890
412	Congenital Myasthenic Syndrome Panel	\$590-5,310	668	<i>NR2E3</i> Sequencing	\$ 540
407	<i>CHAT</i> Sequencing	\$ 960	684	<i>PDE6A</i> Sequencing	\$1,190
401	<i>CHRNA1</i> Gene Sequencing	\$ 680	685	<i>PDE6B</i> Sequencing	\$1,190
402	<i>CHRNB1</i> Gene Sequencing	\$ 680	669	<i>PRPF3</i> Sequencing	\$190-890
403	<i>CHRND</i> Gene Sequencing	\$ 770	667	<i>PRPF8</i> Sequencing	\$190-1,990
404	<i>CHRNE</i> Gene Sequencing	\$ 730	666	<i>PRPF31</i> Sequencing	\$ 740
409	<i>COLQ</i> Sequencing	\$ 920	663	<i>PRPH2</i> Sequencing	\$ 420
406	<i>MUSK</i> Sequencing	\$ 920	661	<i>RHO</i> Sequencing	\$ 540
			662	<i>RP1</i> Sequencing	\$ 1,140
			682	<i>RPE</i> Sequencing	\$ 840
	Cystic Fibrosis		683	<i>TULP1</i> Sequencing	\$ 640
150	<i>CFTR</i> Sequencing + 5T/TG Tract Analysis	\$1,290			
				Fabry Disease	
			471	<i>GLA</i> Sequencing	\$ 490
	Costello Syndrome				
140	<i>HRAS</i> Sequencing	\$190 - 450		Familial Hemophagocytic Lymphohistiocytosis	
141	<i>HRAS</i> Sequencing – Tier 1 only	\$ 190	215	Familial Hemophagocytic Lymphohistiocytosis (FHL) Panel	\$440-3,200
142	<i>HRAS</i> Sequencing – Tier 2 only	\$ 260	216	<i>PRF1</i> Sequencing	\$ 540
			217	<i>STX11</i> Sequencing	\$ 440
			214	<i>STXBP2</i> Sequencing	\$1,060
	Danon Disease		218	<i>UNC13D</i> Sequencing – Tier 1	\$ 840
418	<i>LAMP2</i> Sequencing	\$ 620	219	<i>UNC13D</i> Sequencing – Tier 2	\$ 890
				Familial Limb Girdle Myasthenic Syndrome	
	Distal Arthrogyrosis		408	<i>AGRN</i> Sequencing	\$1,780
337	<i>MYH3</i> Gene Sequencing	\$ 1760	414	<i>DOK7-AGRN</i> Sequencing	\$540-2130
333	<i>TNNI2</i> Gene Sequencing	\$ 530			
334	<i>TNNT3</i> Gene Sequencing	\$ 820			
				Fanconi-Bickel Syndrome (GSD, Type XI)	
	Distal Myopathy		170	<i>SLC2A2</i> Sequencing	\$ 650
336	<i>MATR</i> Gene Sequencing	\$ 820			
	Dynactin-Related Disorders			Farber Lipogranulomatosis	
382	<i>DCTN1</i> Sequencing	\$1,690	481	<i>ASAH1</i> Sequencing	\$ 820
	Dystonia				
160	<i>DRD2</i> Sequencing	\$ 540		Fatty Acid Oxidation Disorders	
161	<i>GCH1</i> Sequencing	\$ 440	180	<i>ACADM</i> Sequencing	\$ 540
162	<i>PNKD</i> Sequencing	\$190 - 540	183	<i>ACADS</i> Sequencing	\$ 540
163	<i>PNKD</i> Sequencing - Tier 1 only	\$ 190	184	<i>ACADVL</i> Sequencing	\$ 740
164	<i>PNKD</i> Sequencing - Tier 2 only	\$ 350			
165	<i>SGCE</i> Sequencing	\$ 590			
166	<i>TOR1A</i> Sequencing	\$190 - 440			
167	<i>TOR1A</i> Sequencing - Tier 1 only	\$ 190	201	Galactosemia	
168	<i>TOR1A</i> Sequencing – Tier 2 only	\$ 290		<i>GALT</i> Sequencing	\$ 440
	Eye Diseases			Glutaric Acidemia I	
686	<i>CNGA1</i> Sequencing	\$ 690	210	<i>GCDH</i> Sequencing	\$ 540
687	<i>CNGB1</i> Sequencing	\$1,590			
641	<i>COL18A1</i> Sequencing	\$1,990			
681	<i>CRB1</i> Sequencing	\$1,040			
665	<i>CRX</i> Sequencing	\$ 440			

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			198	ACTN2 Sequencing	\$1,090
			196	CSRP3 Sequencing	\$ 470
	Glycogen Storage Disease		172	MYH7 Sequencing	\$ 1640
220	GYS2 Sequencing (Type 0)	\$ 940	175	MYL2 Sequencing	\$ 480
221	G6PC Sequencing (Type Ia)	\$ 440	176	MYL3 Sequencing	\$ 450
222	SLC37A4 Sequencing (Type Ib)	\$ 540	178	TNNI3 Sequencing	\$ 570
223	GAA Sequencing (Type II)	\$ 990	179	TNNT2 Sequencing	\$ 820
224	AGL Sequencing (Type III)	\$1,790	177	TPM1 Sequencing	\$ 620
225	GBE1 Sequencing (Type IV)	\$ 990	144	VCL Sequencing	\$1,170
418	LAMP2 Sequencing (Type IIb)	\$ 620			
226	PYGM Sequencing (Type V)	\$340 - 990			
227	PYGM Sequencing (Type V) – Tier 1 only	\$ 340		Inclusion Body Myopathy	
228	PYGM Sequencing (Type V) – Tier 2 only	\$ 790	367	GNE Sequencing	\$ 740
229	PYGL Sequencing (Type VI)	\$1,140	361	MYH2 Sequencing	\$1,690
230	PFKM Sequencing (Type VII)	\$1,090			
231	PHKA1 Sequencing (Type IX)	\$1,690			
232	PHKA2 Sequencing (Type IX)	\$490-1,690		Isovaleric Acidemia	
233	PHKB Sequencing (Type IX)	\$1,690	250	IVD Sequencing	\$ 590
235	PHKG2 Sequencing (Type IX)	\$ 490			
				Joubert Syndrome	
			294	Joubert Syndrome Panel	\$290-8,270
	Glycyl tRNA Synthetase-Related Disorders		266	AHI1 Sequencing	\$320
383	GARS Sequencing	\$ 940	277	ARL13B Sequencing	\$ 640
			278	CC2D2A Sequencing	\$1,590
			267	CEP290 Sequencing	2,190
	GM1 Gangliosidosis/Morquio Syndrome Type B		279	INPP5E Sequencing	\$ 690
477	GLB1 Sequencing	\$ 840	274	TMEM67/MKS3 Sequencing	\$1,290
			275	NPHP1 Deletion Testing	\$ 290
			651	NPHP1 Sequencing	\$1,090
	Heat Shock 22/27 kDa Protein-Related Disorders		276	RPGRIP1L Sequencing	\$1,190
387	HSPB1 Sequencing	\$ 390	291	TMEM216 Sequencing	\$ 460
389	HSPB8 Sequencing	\$ 440			
				Ciliopathy	
			279	INPP5E Sequencing	\$ 690
				Krabbe Disease	
	Holoprosencephaly		631	GALC Sequencing	\$ 830
580	Holoprosencephaly Sequential (SHH, ZIC2, SIX3, TGIF1, GLI2)	\$390-3,290	632	GALC Deletion Test	\$ 190
586	GLI2 Sequencing	\$ 990			
585	PTCH1 Sequencing	\$1,090		Leber Congenital Amaurosis	
581	SHH Sequencing	\$ 470	681	CRB1 Sequencing	\$1,040
583	SIX3 Sequencing	\$ 440	665	CRX Sequencing	\$ 440
584	TGIF1 Sequencing	\$ 390			
582	ZIC2 Sequencing	\$ 540		Limb Girdle Muscular Dystrophy	
			469	ANO5/TMEM16E Sequencing	\$1,090
				Lissencephaly	
	3-Hydroxy-3-Methylglutaryl-CoA Lyase Deficiency		504	TUBA1A Sequencing	\$ 440
240	HMGCL Sequencing	\$ 590	502	ARX Sequencing	\$ 590
			506	RELN Sequencing	\$2,590
	Hypertrophic Cardiomyopathy		505	YWHAE Sequencing	\$ 520
190	Sequential Testing of Eight Sarcomeric Genes	\$5,580			
191	MYBPC3-MYH7 Sequencing	\$2,630		Loeys-Dietz Syndrome	
193	MYL2-MYL3-ACTC1 Sequencing	\$1,240	396	TGFBR1 Sequencing	\$ 620
192	TNNT2-TNNI3-TPM1 Sequencing	\$1,710	397	TGFBR2 Sequencing	\$ 710
174	ACTC1 Sequencing	\$ 530			

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	Lujan Syndrome				
398	MED12 Sequencing	\$1,960		Mucopolidosis	
	Lynch Syndrome		316	GNPTAB Sequencing	\$ 1,190
701	MLH1 Sequencing	\$1,290	317	GNPTG Sequencing	\$ 470
702	MSH2 Sequencing	\$1,090			
703	MSH6 Sequencing	\$1,050			
704	PMS1 Sequencing	\$ 940		Mucopolysaccharidosis	
	Lymphedema		452	Type I – IDUA Sequencing	\$ 840
280	FOXC2 Sequencing	\$ 390	453	Type II – IDS Sequencing	\$ 670
281	FLT4 Sequencing – Tier 1 only	\$ 540	459	Type IIIA – SGSH Sequencing	\$ 710
282	FLT4 Sequencing – Tier 2 only	\$ 950	484	Type IIIB – NAGLU Sequencing	\$ 740
283	SOX18 Sequencing	\$ 490	458	Type IIIC – HGSNAT Sequencing	\$ 960
	Lysosomal Storage Disorders		456	Type IIID – GNS Sequencing	\$ 880
477	GLB1 Sequencing	\$ 840	455	Type IVA – GALNS Sequencing	\$ 840
476	HEXB Sequencing – Tier 1 only	\$ 840	454	Type VI – ARSB Sequencing	\$ 620
	Malignant Hyperthermia /Central Core Disease		457	Type VII – GUSB Sequencing	\$ 660
570	Sequential RYR1 Sequencing	\$740–3,990	485	Type IX – HYAL1 Sequencing	\$ 540
571	RYR1 Sequencing – Tier 1 only	\$ 740			
572	RYR1 Sequencing – Tier 2 only	\$3,790			
	Marfan Syndrome			Multiminicore, Rigid Spine Muscular Dystrophy	
394	FBN1 Sequencing	\$2,190	330	SEPN1 Sequencing	\$ 690
			357	TPM3 Sequencing	\$ 570
	Meckel Gruber Syndrome				
295	Meckel-Gruber Syndrome Sequencing Panel	\$840-6,035		Multiple Carboxylase Deficiency	
278	CC2D2A Sequencing	\$1,590	520	BTD Sequencing	\$ 540
301	MKS1 Sequencing	\$290 - 840	521	HLCS Sequencing	\$ 740
302	MKS1 Sequencing – Tier 1 only	\$ 290			
303	MKS1 Sequencing – Tier 2 only	\$ 790		Multiple Pterygium Syndrome	
276	RPGRIP1L Sequencing	\$1,190	415	CHRNA1, CHRND, CHRNG Sequencing	\$680-1890
274	TMEM67/MKS3 Sequencing	\$1,290	405	CHRNG Sequencing	\$ 730
	Mental Retardation			Multiple Sulfatase Deficiency	
561	CUL4B Sequencing	\$1,090	512	SUMF1 Sequencing	\$ 590
562	SLC9A6 Sequencing	\$ 790			
134	VPS13B Sequencing	\$2,790		Muscular Dystrophy	
				Dystroglycanopathy Sequential (FKRP, FKTN, POMT1, POMT2, POMGnT1, LARGE1)	\$390–4,275
	Metachromatic Leukodystrophy		340		
			341	CAPN3 Sequencing	\$1,090
621	Metachromatic Leukodystrophy Sequencing Panel	\$490-1,140	467	CAV3 Sequencing	\$ 390
511	PSAP Sequencing	\$ 790	384	DAG1 Sequencing	\$ 640
			465	DOK7 Sequencing	\$ 540
	Methylmalonic Acidemia		342	DYSF Sequencing	\$2,490
310	Sequential Gene Sequencing	\$790-1550	344	FKRP Sequencing	\$ 390
311	MUT Sequencing	\$ 790	343	FKTN /FCMD Sequencing	\$ 590
312	MMAA Sequencing	\$ 490	244	ITGA7 Sequencing	\$1,240
313	MMAB Sequencing	\$ 540	345	LAMA2 Sequencing	\$2,990
	3-Methylcrotonyl-CoA Carboxylase Deficiency		245	LAMA2: Mexican Exon 55 Mutation	\$ 190
320	Sequential Gene Sequencing	\$840-1,770	346	LARGE Sequencing	\$ 890
321	MCCC2 Sequencing	\$ 840	347	LMNA Sequencing	\$ 590
322	MCCC1 Sequencing	\$ 990	366	MYOT/TTID Sequencing	\$ 190-590
			355	NEB Sequencing	\$3245-6490
	3-Methylglutaconic Aciduria Type I		356	NEB Exon 55 Deletion Test	\$ 200
323	AUH Sequencing	\$ 690			
			348	POMT1 Sequencing	\$ 890

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			349	POMT2 Sequencing	\$ 990
351	POMGNT1 Sequencing	\$ 640		Niemann-Pick Disease	
330	SEPN1 Sequencing	\$ 690	473	NPC1 Sequencing	\$1,240
352	SGCA Sequencing	\$ 490	474	NPC2 Sequencing	\$ 490
353	SGCB Sequencing	\$ 440			
338	SGCG Sequencing	\$ 570		Noonan Syndrome/LEOPARD Syndrome	
246	SYNE1 Sequencing	\$6,490	370	Sequential (PTPN11, SOS1, RAF1, KRAS)	\$390 - 1,990
247	SYNE1 Seq. - French Canadian Mutation Panel	\$ 550	371	Sequential – Tier 1 only	\$ 390
468	TCAP Sequencing	\$ 390	372	Sequential – Tier 2 only	\$1,990
385	TTN Sequencing	\$ 1290	373	Noonan PTPN11 Sequencing	\$ 390 - 890
			371	Noonan PTPN11 Sequencing – Tier 1 only	\$ 390
	Myofibrillar Myopathy		374	Noonan PTPN11 Sequencing – Tier 2 only	\$ 690
362	CRYAB Sequencing	\$ 370	375	RAF1 Sequencing	\$ 370
363	DES Sequencing	\$ 670	376	SOS1 Sequencing	\$590 - 1,190
364	FLNC Sequencing	\$2,490	377	LEOPARD PTPN11 Sequencing	\$370 - 890
365	LDB3 (ZASP) Sequencing	\$ 940	378	KRAS Sequencing	\$ 440
			115	NF1 Sequencing	\$540-2490
			379	NRAS Sequencing	\$ 430
	Multiple Sulfatase Deficiency		381	SHOC2 Sequencing	\$ 660
512	SUMF1 Sequencing	\$ 590			
				Pallister-Hall/Greig Cephalopolysyndactyly	
			380	GLI3 Sequencing	\$1,090
	Myotilinopathy				
366	MYOT/TTID Sequencing	\$ 190-590		Palmoplantar Keratoderma	
			208	JUP Sequencing	\$ 730
	Nemaline Myopathy			Pena-Shokeir Syndrome, Type 1	
358	ACTA1 Sequencing	\$ 530	413	DOK7-RAPSN Sequencing Panel	\$540-\$1060
249	CFL2 Sequencing	\$ 420			
		\$3245-6490			
355	NEB Sequencing	\$3245-6490		Phenylalanine Hydroxylase Deficiency (PKU)	
356	NEB Exon 55 Deletion Test	\$ 200	530	PAH Testing	\$ 690
331	TPM2 Sequencing	\$ 590			
357	TPM3 Sequencing	\$ 570		Progeria, Hutchinson-Gilford	
332	TNNT1 Sequencing	\$ 690	347	LMNA Sequencing	\$ 590
	Nephronophthisis				
268	Nephronophthisis Sequencing Panel	\$540-8,670		Propionic Acidemia	
656	GLIS2/NPHP7 Deletion Test	\$ 540	390	Sequential Gene Sequencing	\$740-1,890
652	INVS/NPHP2 Deletion Test	\$ 940	391	PCCA Sequencing	\$1,290
655	IQCB1/NPHP5 Deletion Test	\$ 740	392	PCCB Sequencing	\$ 790
657	NEK8/NPHP9 Sequencing	\$ 840			
275	NPHP1 Deletion Test	\$ 290			
651	NPHP1 Sequencing	\$1,090		Rapsyn-Related Disorders	
653	NPHP3 Sequencing	\$1,320	466	RAPSN Sequencing	\$ 590
654	NPHP4 Sequencing	\$1,350			
				Restrictive Dermopathy, Mandibuloacral Dysplasia	
117	NF1 Sequencing	\$2,490	400	ZMPSTE24 Sequencing	\$ 690
118	NF2 Sequencing	\$ 840			
116	SPRED1 Sequencing	\$ 540			
				Robinow Syndrome, Autosomal Recessive	
			136	ROR2 Sequencing	\$ 740
	Neurofibromatosis-Noonan Syndrome				
115	NF1 Sequencing	\$540-2490		Sandhoff Disease	

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			476	HEXB Sequencing	\$ 840
	Seipin-Related Disorders		433	GP1BA Sequencing	\$ 490
461	BSCL2 Sequencing	\$ 740	434	GP1BB Sequencing	\$ 390
			435	GP9 Sequencing	\$ 340
			436	MASTL Sequencing	\$ 740
	Sialuria		441		\$690-1740
			442	MYH9 Sequencing	
368	GNE, Exon 5 Sequencing	\$ 190	437	MPL Sequencing	\$ 640
			439	RUNX1 Sequencing	\$ 490
			440	WAS Sequencing	\$ 540
	Simpson-Golabi-Behmel Syndrome				
563	GPC3 Sequencing	\$ 590		Tropomyosin 3-Related Myopathies	
			357	TPM3 Sequencing	\$ 570
	Skeletal Disorders			Tyrosinemia	
420	Sequential Test (DLL3, MESP2, LFNG)	\$540-1,290	541	FAH Sequencing	\$ 690
421	DLL3 Sequencing	\$ 590		Tyrosine Hydroxylase Deficiency-Related Disorders	
424	GDF6 Sequencing	\$ 490	159	TH Sequencing	\$ 840
422	LFNG Sequencing	\$ 490			
423	MESP2 Sequencing	\$ 390		Ullrich Congenital Muscular Dystrophy	
			359	COL6A1-COL6A2-COL6A3 Sequencing	\$1290-3890
	Smith-Lemli-Opitz Syndrome		552	Urea Cycle Disorders	
410	DHCR7 Sequencing	\$ 440	553	ARG1 Sequencing	\$ 490
			554	ASS1 Sequencing	\$ 740
			555	CPS1 Sequencing	\$1,890
	Sotos Syndrome		551	NAGS Sequencing	\$ 430
132	NSD1 Sequencing	\$1,490		OTC Sequencing	\$ 540
				Ventricular Tachycardia	
			450	RYR2 Sequencing	\$1,390
	Sphingolipidoses			Vision Disorders	
620	ARSA Sequencing	\$ 490		COL18A1 Sequencing	\$1,990
479	GBA Sequencing	\$ 630	641	White Matter Disorders	
	Spinal Muscular Atrophy with Respiratory Distress Type 1		610	EIF2B1 – EIF2B5 Sequential Sequencing	\$690-2,390
339	IGHMBP2 Sequencing	\$ 910			
				Wilson Disease	
			460	ATP7B Sequencing	\$ 990
	Stuttering			Wolff-Parkinson-White Syndrome	
315	Stuttering Panel (GNPTAB, GNPTG, NAGPA)	\$1,790		PRKAG2 Sequencing	\$ 870
316	GNPTAB Sequencing	\$1,240	199		
317	GNPTG Sequencing	\$ 490		Wolman Disease	
318	NAGPA Sequencing	\$ 690	482	LIPA Sequencing	\$ 620
	Tay-Sachs Disease				
478	GM2A Sequencing	\$ 440		- Other Tests -	
475	HEXA Sequencing	\$ 770	500	DNA Banking	\$ 98
			480	Sex Chromosome Panel	\$ 90
	Thrombocytopenia		490	DNA Identity Panel	\$ 90
430	ADAMTS13 Sequencing	\$1,490		Gender Analysis, Zygosity Testing, Specimen Matching	
431	GATA1 Sequencing	\$ 390	510	Genome Polymorphism Scan	\$ 590
432	Bernard-Soulier Sequential(GP1BA, GP1BB, GP9)	\$490 - 890			



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